

Supplementary Data

Facile and scalable fabrication of three-dimensional Cu(OH)₂ nanoporous nanorods for solid-state supercapacitors†

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Calculation methods

1. Three-electrode configuration

The areal capacitance C_a (F cm⁻²) and gravimetric capacitance C_g (F g⁻¹) of the Cu(OH)₂ could be calculated from the corresponding cyclic voltammetry curve by the following equations:

$$C_a = \frac{\int I(V)dV}{2\nu S\Delta V} \quad (1)$$

$$C_g = \frac{\int I(V)dV}{2\nu m\Delta V} \quad (2)$$

where $I(V)$ (A) is the response current, V (V) is the potential vs. SCE, ν (V s⁻¹) is the scan rate, S (cm²) is the effective area of the electrode, m (g) is the mass of the active material on the electrode, ΔV (V) is the working potential.

C_a (F cm⁻²) and C_g (F g⁻¹) could also be calculated from the corresponding galvanostatic discharging curve using the following equations:

$$C_a = \frac{I\Delta t}{S\Delta V} \quad (3)$$

$$C_g = \frac{I\Delta t}{m\Delta V} \quad (4)$$

where I (A) is the discharging current, Δt (s) is the discharging time, S (cm²) is the effective area of the electrode, m (g) is the mass of the active material on the electrode, ΔV (V) is the working

potential.

The equivalent series resistance (R_{ESR}) (Ω) could be calculated by:

$$R_{ESR} = \frac{V_{drop}}{2I} \quad (5)$$

where V_{drop} (V) is the abrupt voltage drop at the beginning of the discharging curve, I (A) is the corresponding current.

2. Two-electrode system (asymmetric supercapacitor, denoted as ASC)

The specific capacitance C_{ASC} ($F\ g^{-1}$) of the ASC could be calculated from the corresponding galvanostatic discharging curve according to the following equation:

$$C_{ASC} = \frac{I\Delta t}{M\Delta V} \quad (6)$$

where I (A) is the discharging current, Δt (s) is the discharging time, ΔV (V) is the potential window during the discharging process, M (g) is the total mass of active materials in the ASC.

The energy density E ($mWh\ cm^{-3}$) and average power density P ($mW\ cm^{-3}$) could be calculated as:

$$E = \frac{\int V(t)dtI}{3.6Vol} \quad (7)$$

$$P = \frac{3600 E}{t} \quad (8)$$

where $V(t)$ (V) is the potential of the ASC, t (s) is the corresponding discharging time, I (A) is the discharging current, Vol (cm^3) is the volume of the ASC including the volumes of two Cu foil current collectors and the gel electrolyte.

Supplementary Figures

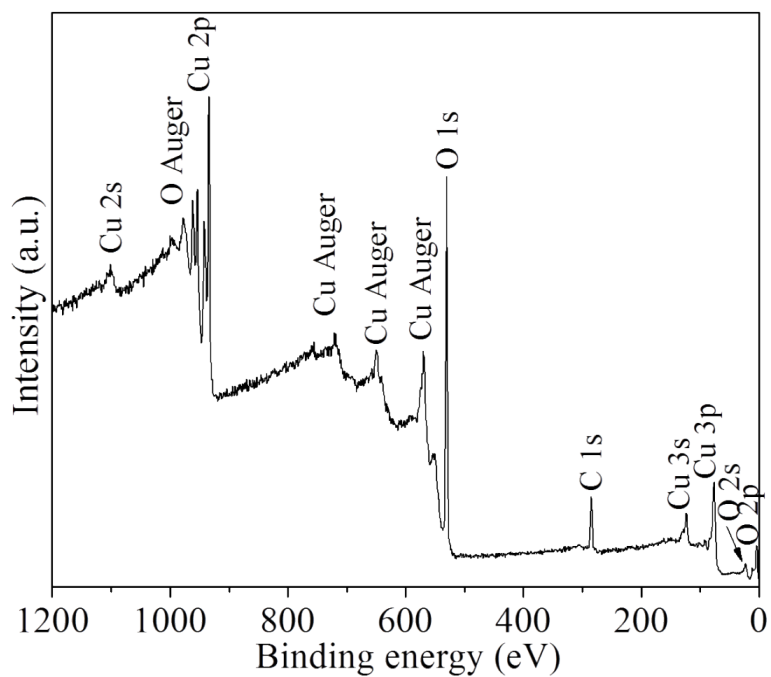


Fig. S1 Survey XPS spectrum of 3-D $\text{Cu}(\text{OH})_2$.

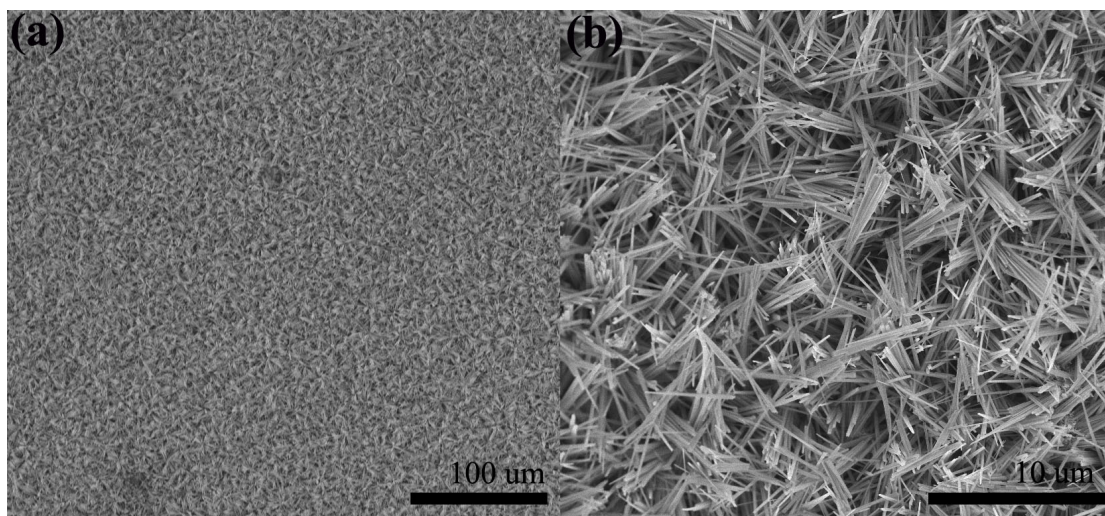


Fig. S2 (a,b) Low-magnification SEM images of 3-D Cu(OH)₂.

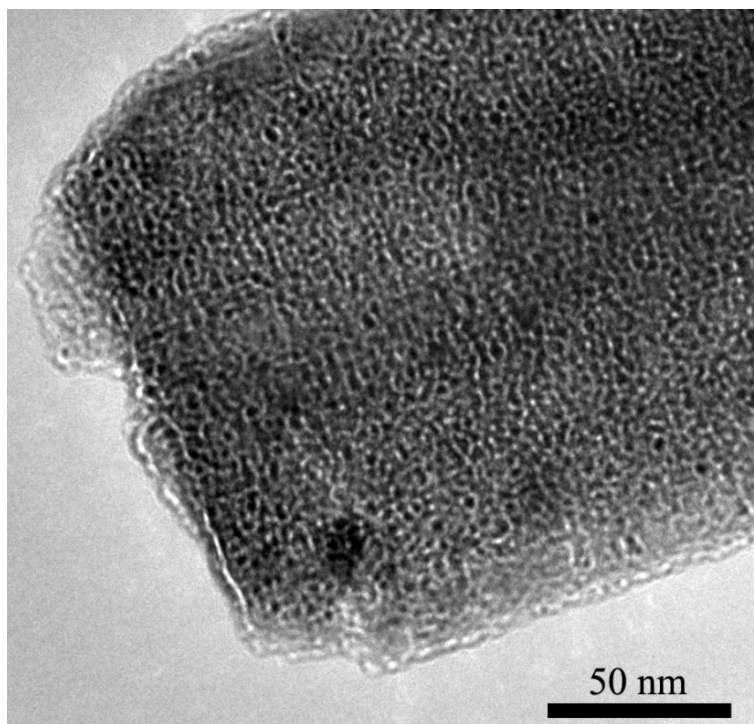


Fig. S3 Magnified TEM image of 3-D Cu(OH)₂ from Fig. 3d.

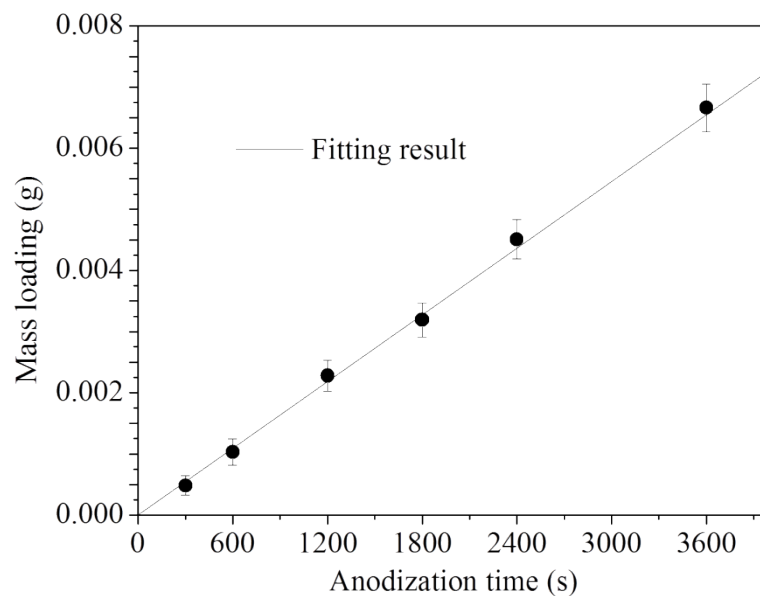


Fig. S4 Mass loading of $\text{Cu}(\text{OH})_2$ on the $2 \times 3 \text{ cm}^2$ $\text{Cu}(\text{OH})_2/\text{Cu}$ slice ($2 \times 2 \text{ cm}^2$ covered by $\text{Cu}(\text{OH})_2$) with respect to different anodization time.

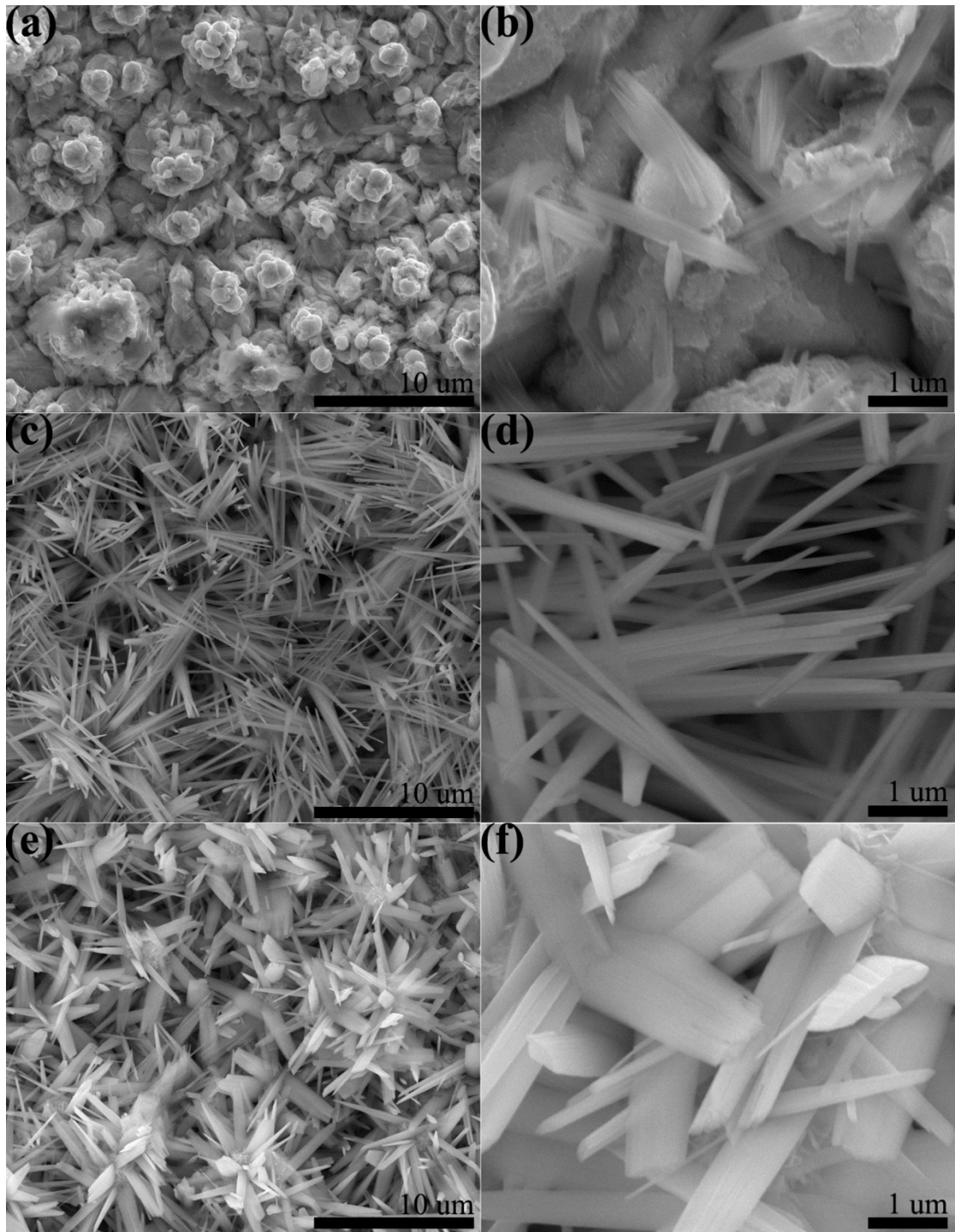


Fig. S5 SEM images of 3-D Cu(OH)₂ with respect to the different anodization time: (a, b) 100 s; (c, d) 1800 s; (e, f) 3600 s.

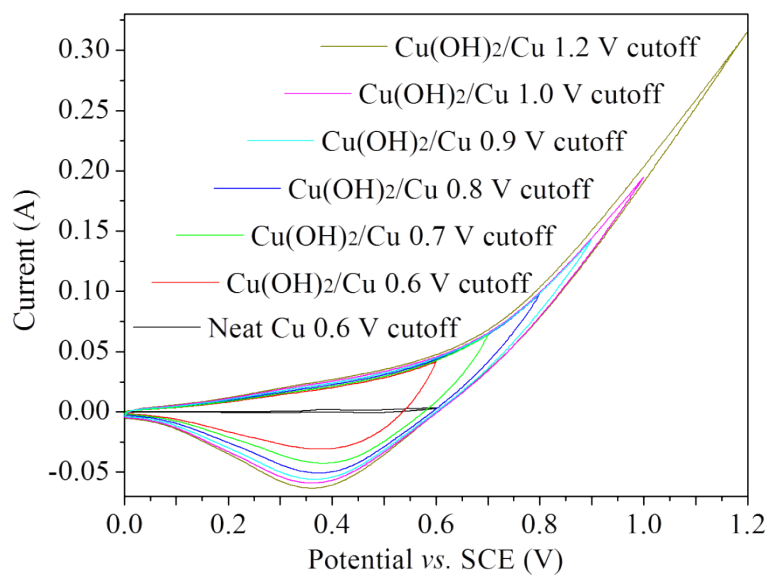


Fig. S6 CV curves of the neat Cu slice with a potential cutoff of 0.6 V at 50 mV s⁻¹ and the Cu(OH)₂/Cu slice with potential cutoffs from 0.6 to 1.2 V at 50 mV s⁻¹.

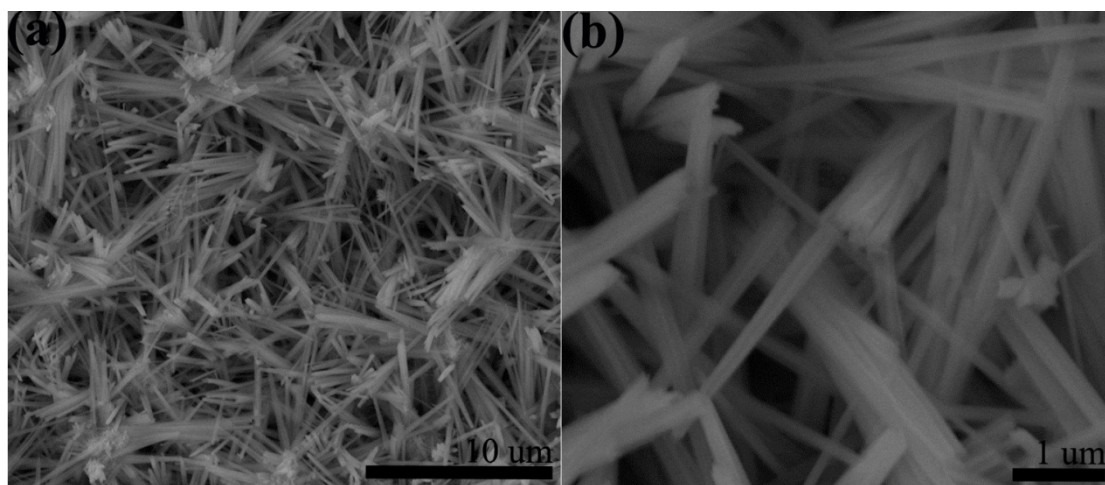


Fig. S7 (a,b) SEM images of 3-D Cu(OH)₂ after 5000 cycles.

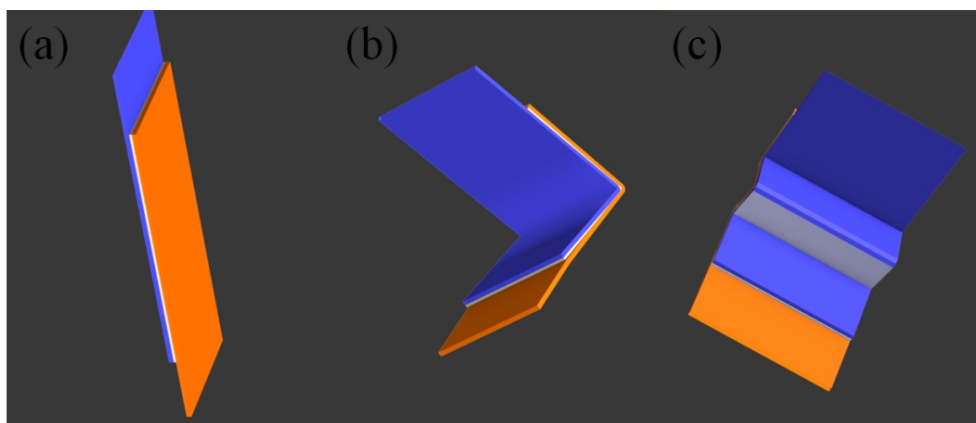


Fig. S8 Graphic illustrations of the Cu(OH)₂//AC solid-state ASC under (a) normal state, (b) bended by one fold and (c) bended by three folds.

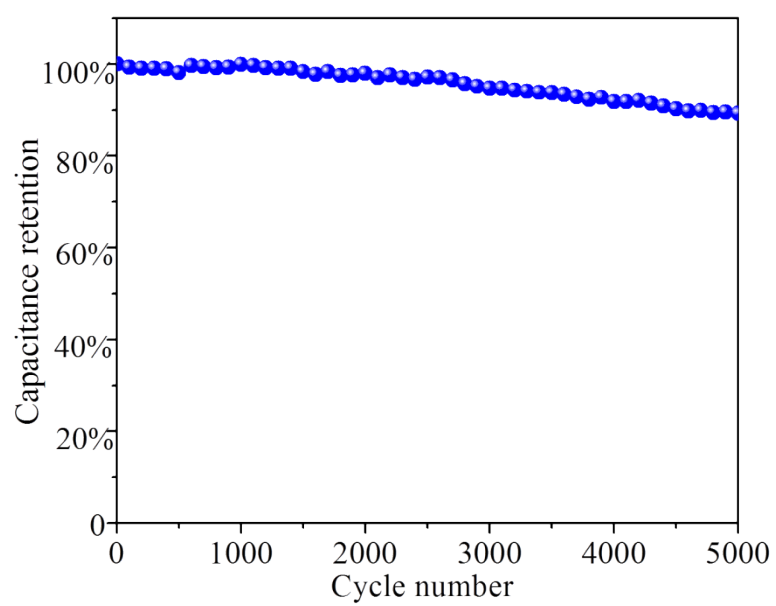


Fig. S9 Cycling performance of Cu(OH)₂//AC solid-state ASC at 10 A g⁻¹.

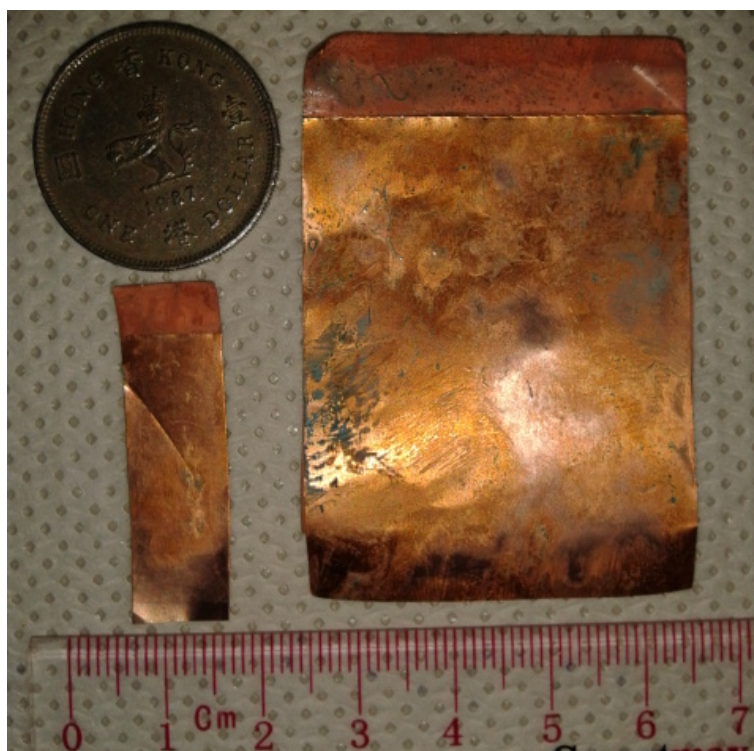


Fig. S10 Photographs of $1 \times 4 \text{ cm}^2$ (left) and $4 \times 6 \text{ cm}^2$ (right) $\text{Cu}(\text{OH})_2//\text{AC}$ solid-state ASCs.

Video S1 Demonstration of our tandem device (consisting of two $4 \times 6 \text{ cm}^2$ solid-state ASCs) powering 26 LEDs.